AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application.

What is claimed is:

 (Currently Amended) An injectable radiological composition for x-ray visualization during radiological examinations, the composition comprising a pharmaceutically acceptable vehicle and a mixture of at least one monomer and at least one dimer, the monomer corresponding to Formula I and the dimer corresponding to Formula II

$$\begin{array}{c} A_1 \\ B_1 \\ B_1 \\ E_2 \\ X \\ E_3 \\ \end{array}$$
 Formula (I)

wherein, with regard to Formula I:

A₁[,] and B₁, and D₄ are independently -CON(R₃)R₁; or

 \underline{D}_1 is $-N(R)C(O)R_2$;

 $A_{z_1}A_{z_2}-B_{z_3}-\text{and}-D_{z_2}\text{ are independently-CON(R)}R_1-\text{or-N(R)C(O)}R_{z_2}\text{ provided, however, at least one of }A_{z_2}\text{ and }A_{z_3}\text{ is-CONH}_{z_1}$

 $E_2 \text{ and } - E_3 \text{ are independently selected from the group consisting of -CON(R)}, -N(R)C(O) \\ \text{and -N(COR_2)};$

each R and R₂ is independently H, or a linear or branched (C₁-C₈) alkyl residue, optionally substituted by one or more hydroxy, alkoxy or hydroxyalkoxy groups or combinations thereof, or a member of a (C₄ - C₇) cyclic residue, said-cyclic residue being optionally interrupted by O - S or NR₄, and/or optionally substituted with one or more hydroxy, alkoxy or

hydroxyalkoxy groups or combinations thereof, the cyclic residue comprising R, the nitrogen atom to which it is bonded and another moiety, that moiety being (i) –C(O)R₂ when A₁, A₂, A₃, B₁, D₁ or D₂ is –N(R)C(O)R₂ or (ii) R₁ when A₂, A₃, B₃, or D₂ is –CON(R)R₁:

each R_1 is independently (i) hydrogen, \underline{or} (ii) a linear or branched (C_1-C_8) alkyl residue, optionally substituted with one or more hydroxy, alkoxy, hydroxyalkoxy groups or combinations thereof or by $-NRC(O)R_4$ -or $-C(O)N(R)R_4$, (iii) the residue of a carbohydrate, or (iv) a member of a $-(C_3-C_2)$ cyclic residue, said cyclic residue being optionally interrupted by $-O_-$, $-S_-$ or $-NR_4$, and/or optionally substituted by one or more hydroxy, alkoxy or hydroxyalkoxy groups or combinations thereof, the cyclic residue comprising R_4 , the nitrogen atom to which it is bonded and another molety, that molety being (a) R when A_2 , A_3 , B_4 , or $-C_2$ or $-C_3$ or $-C_4$ or $-C_4$, or $-C_4$, $-C_4$, -

each R₂-is independently (i) a linear or branched (C₄—C₆) alkyl-residue, optionally substituted with one or more hydroxy, alkexy or hydroxyalkexy-groups, or combinations thereof or (ii) a member of a (C₄—C₇) syclic residue, said cyclic residue being optionally interrupted by O₇—S or NR₄—and/or optionally substituted with one or more hydroxy, alkexy or hydroxyalkexy groups or combinations thereof, the cyclic residue comprising R₂, R, the nitrogen atom to which R is bended and the carbonyl mojety to which R₂ is bended:

each R_3 is independently linear or branched (C_1 – C_8) alkyl residue, optionally substituted with one or more hydroxy, alkoxy or hydroxyalkoxy groups or combinations thereof, or taken together with R_1 -and the nitrogen atom to which R_3 -and R_1 are bended, form a (C_3 – C_2)-cyclic residue, said cyclic residue being optionally interrupted by $-O_1$, $-S_2$ -or $-NR_4$, and/or optionally substituted by one or more hydroxy, alkoxy or hydroxyalkoxy groups or combinations thereof;

each- R_4 -is independently hydrogen or a linear or branched $(G_4 - G_6)$ alkyl-residue, optionally substituted with one or more hydroxy, alkoxy, hydroxyalkoxy groups or combinations thereof; and

X is a bond or a linear or branched $(C_1 - C_6)$ alkylene chain which is optionally substituted by up to six hydroxy groups, said alkylene chain being optionally interrupted by $-O_1 - O_2$.

S., $-NR_1 - or - N(R)C(O)_2$ groups.

and wherein with regard to Formula II:

A₂ and A₃ are -CONH₂;

B₃ and D₂ are -CON(R)R₁;

E₂ and E₃ are independently selected from the group consisting of -CON(R)-, -N(R)C(O)- and -N(COR₂)-;

each R is independently H, or a linear or branched (C₁-C₈) alkyl residue, optionally

substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof, each R_1 is independently (i) hydrogen, (ii) a linear or branched (C_1 - C_8) alkyl residue, optionally substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof, or (iii) the residue of a carbohydrate:

or R and R_1 are each members of a (C_3-C_7) cyclic residue further comprising the nitrogen atom to which each of R and R_1 is bonded, said cyclic residue being optionally interrupted by $-O_-$, $-S_-$ or $-NR_2$, and/or optionally substituted by one or more hydroxy, alkoxy or hydroxyalkoxy groups or combinations thereof;

 $\frac{\text{each } R_2 \text{ is independently a linear or branched } (C_1\text{-}C_8) \text{ alkyl residue, optionally}}{\text{substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof;}}\\ \frac{\text{each } R_4 \text{ is independently hydrogen or a linear branched } (C_1\text{-}C_8) \text{ alkyl residue, optionally}}{\text{substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof;}}\\ \text{and}$

X is a bond or a linear or branched (C_1-C_8) alkylene chain which is optionally substituted by up to six hydroxyl groups, said alkylene chain being optionally interrupted by -O-, -S-, -NR₄- or -N(R)C(O)- groups.

- $\label{eq:composition} 2. \mbox{ (Currently Amended) The composition of claim 1 wherein A_2 and A_3 are independently C(O)NH$_2$ with regard to Formula I, R_1 is H or methyl. }$
 - 3. (Original) The composition of claim 1 wherein X is methylene.
- 4. (Currently Amended) The composition of claim 1 wherein A₄-and-B₄-are-C(O)N(R₃)R₄, and each R₃-and R₄-of A₄-and B₄-are as defined in claim 1 with regard to Formula L:

 A_1 and B_1 are -CON(R_3) R_1 ;

D₁ is -N(R)C(O)R₂:

each R and R₂ is independently H, methyl, hydroxymethyl, ethyl, hydroxyethyl, propyl, hydroxypropyl, 2-methoxyethyl, 1-methoxy-2-hydroxypropyl or dihydroxypropyl;

each R₁ is independently H or methyl;

each R₃ is independently methyl, hydroxymethyl, ethyl, hydroxyethyl, propyl,

hydroxypropyl or dihydroxypropyl;

and wherein with regard to Formula II:

A₂ and A₃ are -CONH₂;

B₃ and D₂ are -CON(R)R₁:

E₂ and E₃ are independently selected from the group consisting of -CON(R)-, -N(R)C(O)and -N(COR₂)-;

each R is independently H, or a linear or branched $(C_1 - C_8)$ alkyl residue, optionally substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof, each R_1 is independently (i) hydrogen, (ii) a linear or branched $(C_1 - C_8)$ alkyl residue, optionally substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof or by -NRC(O)R₁ or -C(O)N(R)R₁, or (iii) the residue of a carbohydrate;

or R and R_1 are each members of a $(C_3$ - $C_7)$ cyclic residue further comprising the nitrogen atom to which each of R and R_1 is bonded, said cyclic residue being optionally interrupted by -O-, -S- or -NR $_2$ -, and/or optionally substituted by one or more hydroxy, alkoxy or hydroxyalkoxy groups or combinations thereof:

 $\frac{\text{each } R_2 \text{ is independently a linear or branched } (C_1\text{-}C_8) \text{ alkyl residue, optionally}}{\text{substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof,}}\\ \frac{\text{each } R_4 \text{ is independently hydrogen or a linear branched } (C_1\text{-}C_8) \text{ alkyl residue, optionally}}{\text{substituted by one or more hydroxyl, alkoxy or hydroxyalkoxy groups or combinations thereof,}}\\ \text{and}$

X is a bond or a linear or branched (C_1-C_8) alkylene chain which is optionally substituted by up to six hydroxyl groups, said alkylene chain being optionally interrupted by -O-, -S-, -NR₄- or -N(R)C(O)- groups.

- 5. (Cancelled)
- (Currently Amended) The composition of claim 1 wherein A₁ and B₁ are -CONHR₃ wherein each R₂ of A₁-and B₁ is independently methyl, hydroxymethyl, ethyl, hydroxyethyl, propyl, hydroxypropyl, or dihydroxypropyl.
 - 7. (Cancelled)
- (Withdrawn Currently Amended) The composition of claim 1 wherein A₁-and-B₂-are
 -CONR₄R₃ wherein each R₁ and R₃ of A₁ and B₁ is independently methyl, hydroxymethyl, ethyl, hydroxyethyl, propyl, hydroxypropyl, or dihydroxypropyl.
 - 9. (Cancelled)

- (Currently Amended) The composition of claim 1 wherein D₁-is-N(R)C(O)R₂, and the R and R₂ substituents of D₁ are independently methyl, hydroxymethyl, ethyl, hydroxyethyl, propyl, hydroxypropyl, 1-methoxy-2-hydroxypropyl, or dihydroxypropyl.
- (Currently Amended) The composition of claim 10 wherein A₁ and B₁ are -CONHR₃ wherein each R₂ of A₄-and B₂ is independently methyl, hydroxymethyl, ethyl, hydroxygropyl, or dihydroxypropyl.
- 12. (Withdrawn Currently Amended) The composition of claim 10 wherein A₄-and-B₄ are—CONR₄R₃-wherein each R₁ and R₃ of A₁ and B₁ is independently methyl, hydroxymethyl, ethyl, hydroxyethyl, propyl, hydroxypropyl, or dihydroxypropyl.
- 13. (Currently Amended) The composition of claim $2\underline{1}$ wherein at least one of A_{ij} . By and B_{ij} is $CONR_{ij}R_{ij}$ wherein R_{1} is hydrogen.
- (Currently Amended) The composition of claim £ 1 wherein ene of A₁, B₂ and D₄ is N(R)C(O)R₂ and R and R₂ are as defined in claim 1 B₃ and D₂ are -CONHR.
- (Original) The composition of claim 1 wherein the monomer is selected from the group consisting of iomeprol, iopromide, ioversol, iohexol, iopentol, iopamidol and iobitridol.
 - 16. (Original) The composition of claim 1 wherein the dimer is josmin.
- 17. (Original) The composition of claim 1 wherein the monomer is selected from the group consisting of ioversol, iohexol, and iopamidol, and the dimer is iosmin.
- (Original) The composition of claim 1 wherein the monomer is ioversol and the dimer is iosmin.
- 19. (Original) The composition of claim 1 wherein the composition further comprises pharmaceutically acceptable radiological vehicles selected from the group consisting of aqueous buffer solutions, sterile water for injection, physiologic saline, balanced ionic solutions, a chelating agent, and other non-radioactive additives comprising excipients and anticlotting

agents.

- 20. (Original) The composition of claim 19 wherein said aqueous buffer solutions comprise tris(hydroxyethyl)amino methane and salts thereof, phosphate, citrate and bicarbonates; wherein said balanced ionic solutions comprise chlorides and bicarbonates of cations selected from the group consisting of Ca, Na, K, and Mg, and other halides, carbonates, sulphates, phosphates of Na, K, Mg and Ca; wherein said chelating agents consist of H₄EDTA, EDTACaNa₂ and calcium monosodium DTPA-BMEA; wherein said excipient is glycerol, polyethylene glycol or dextran; and wherein said anticlotting agent is heparin or hirudin.
- 21. (Withdrawn) The composition of claim 1 wherein the composition further comprises a contrast agent other than the monomer and the dimer.
- 22. (Withdrawn) The composition of claim 21 wherein said other contrast agent is selected from the group consisting of other X-ray contrast agents, magnetic resonance imaging agents, radionuclide imaging agents, ultrasound imaging agents and optical imaging agents.
- 23. (Withdrawn) A method of diagnostic imaging, the method comprising administering to an individual a composition of claim 1, and carrying out an imaging procedure on such individual.
- 24. (Withdrawn) The method of claim 23 wherein said composition comprises a monomer selected from the group consisting of ioversol, iohexol and iopamidol, and the dimer is iosimenol.
- (Withdrawn) The method of claim 23 wherein said composition comprises a mixture of ioversol, and iosimenol.
- 26. (Withdrawn) A method of diagnostic imaging, the method comprising administering to an individual a composition of claim 22, and carrying out an imaging procedure on such individual.